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RESULT 2
US-09-981-353-54
; Sequence 54, Application US/09981353
 Patent No. US20020160382A1
 GENERAL INFORMATION:
  APPLICANT: Lasek, Amy W.
  APPLICANT: Jones, David A.
  TITLE OF INVENTION: GENES EXPRESSED IN COLON CANCER
  FILE REFERENCE: PA-0038 US
  CURRENT APPLICATION NUMBER: US/09/981,353
  CURRENT FILING DATE: 2001-10-11
  NUMBER OF SEQ ID NOS: 194
  SOFTWARE: PERL Program
 SEQ ID NO (54)
   LENGTH: 917
   TYPE: PRT
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   OTHER INFORMATION: Incyte ID No. US20020160382A1 2771481CD1
US-09-981-353-54
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                           Score 4771; DB 10;
                                            Length 917;
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     05-JUN-2002
                   (first entry)
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     Human calcium-activated chloride channel <a href="https://doi.org/10.1001/journal.org/">https://doi.org/10.1001/journal.org/</a>.
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     Nucleic acid library; immune response; asthma; COPD;
KW
     airway hyperresponsiveness; bronchoalveolar manifestation;
     signature sequence; SS; chronic obstructive pulmonary disease;
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     allergic disease; rhinitis; atopic dermatitis; urticaria;
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     autoimmune disease; multiple sclerosis; inflammatory bowel disease;
KW
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     allograft rejection; infectious disease.
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XX
ΡI
     Groot PC, Van Bergenhenegouwen BJ, Van Oosterhout AJM;
XX
DR
     WPI; 2002-241888/29.
XX
PT
     Nucleic acid library comprising genes which are capable of initiation,
     progression and suppression of an immune response, especially an immune
PT
PT
     response observed with airway hyper-responsiveness of asthma -
XX
PS
     Disclosure; Fig 14; 120pp; English.
XX
CC
     The invention relates to a nucleic acid library comprising genes or
CC
     their fragments which are capable of modulating an immune response
CC
     observed with airway hyperresponsiveness and/or bronchoalveolar
CC
     manifestations of asthma. Also included are a method for modulating an
CC
     immune response of an individual comprising modulating a gene comprising
CC
     a nucleic acid at least functionally equivalent to a nucleic acid
     identifiable by a signature sequence (SS) given in the specification such
CC
CC
     as R1-SO-R1-A11, St01-A10, Sv02-1-C11, St01-A12, and R1-SO-R1-B7, a
CC
     substance (for use as a medicament) capable of modulating a gene
CC
     comprising a nucleic acid at least functionally equivalent to a nucleic
CC
     acid identifiable by SS and the use of a proteinaceous substance derived
CC
     from a nucleic acid at least functionally equivalent to a nucleic acid
CC
     identifiable by SS for the production of an antagonist (for use as a
CC
     medicament) against the substance. The antagonist and substance are
CC
     useful for the treatment of an immune response observed with airway
CC
     hyperresponsiveness and/or bronchoalveolar manifestations of asthma.
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The method is useful for modulating the above immune response, where the

CC

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CC
    The substance is useful for treating an immune response, particularly
CC
    asthma, chronic obstructive pulmonary disease (COPD), allergic diseases
CC
    (rhinitis, atopic dermatitis, urticaria), autoimmune diseases (e.g.
CC
    multiple sclerosis), inflammatory bowel disease, allograft rejection and
CC
    infectious disease. The present sequence is a mouse or human
CC
    protein encoded by a signature sequence gene or its homologue/functional
CC
    equivalent.
XX
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                            Mismatches
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Qу
           Db
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Qу
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gene encodes a gene product capable of modulating the immune response.

CC

Db	601 NKDVNSFPSPMIVYAEILQGYVPVLGANVTAFIESQNGHTEVLELLDNGAGADSFKNDGV 660
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Db	901 SVIGSVVIVNFILSTTI 917

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AC
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DT
     20-MAY-2003 (first entry)
XX
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     Amino acid sequence of disease-associated CLCA4 protein.
XX
KW
     Antiinflammatory; Antiasthmatic; Respiratory; Opthalmological;
     Antiallergic; Gastrointestinal; Chest disease;
KW
     Respiratory disease; Bowel disease; Allergic conjunctivitis;
KW
     CLCA4; Human.
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XX
os
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XX
PN
     WO2003005024-A1.
XX
PD
     16-JAN-2003.
XX
PF
     03-JUL-2002; 2002WO-JP06730.
XX
     04-JUL-2001; 2001JP-0203036.
PR
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     (TAKE ) TAKEDA CHEM IND LTD.
XX
     Nakanishi A, Morita S;
PI
XX
     WPI; 2003-210385/20.
DR
XX
PT
     Disease-associated gene CLCA4, its product and antibody, applicable in
PT
     diagnosis and screening drugs for pulmonary and chest diseases
PΤ
     accompanied by inflammation in lung or airway, and respiratory diseases
PT
XX
PS
     Claim 1; Page 63-67; 84pp; Japanese.
XX
CC
     This invention relates to CLCA4, which is applicable in diagnosis
     and screening of drugs for certain diseases and is thought to be
CC
CC
     antiinflammatory, antiasthmatic; opthalmological and antiallergic
CC
     in its action. The CLCA4 gene and its product are applicable in
     diagnosis and screening drugs for pulmonary and chest diseases
CC
CC
     accompanied by inflammation in lung or airway, respiratory diseases
CC
     inflammatory bowel diseases and allergic conjunctivitis. The
CC
     present sequence is the CLCA4 protein. The nucleotide sequence is
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     given in file ABZ59766.
XX
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Db		KIRQRIVCLVLDKSGSMGGKDRLNRMNQAAKHFLLQTVENGSWVGMVHFDSTATIVNKLI 360
Qy		QIKSSDERNTLMAGLPTYPLGGTSICSGIKYAFQVIGELHSQLDGSEVLLLTDGEDNTAS 420
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     05-APR-2000
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                      98US-0093339.
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     30-JUL-1998;
                      98US-0094651.
PR
     04-AUG-1998;
                      98US-0095282.
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PR
     04-AUG-1998;
                     98US-0095285.
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₽R
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                     98US-0096949.
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     18-AUG-1998;
                     98US-0096950.
     18-AUG-1998;
                     98US-0096959.
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     26-AUG-1998;
     26-AUG-1998;
PR
                      98US-0097952.
PR
     26-AUG-1998;
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PR
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                      98US-0097955.
PR
     26-AUG-1998;
                     98US-0097971.
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     26-AUG-1998;
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                      98US-0097986.
     26-AUG-1998;
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PR
PR
     31-AUG-1998;
                     98UŞ-0098525.
PR
     16-SEP-1998;
                      98US-0100634.
PR
     12-JAN-1999;
                      99US-0115565.
XX
PA
      (GETH ) GENENTECH INC.
XX
                Chen J,
PΙ
                          Goddard A, Gurney AL,
                                                   Smith V,
                                                              Watanabe CK;
     Baker K,
ΡI
     Wood WI,
                Yuan J;
XX
     WPI; 2000-072883/06.
DR
DR
     N-PSDB; AAZ65095.
XX
PT
     Membrane-bound proteins and related nucleotide sequences -
XX
PS
     claim 12; Fig 274; 822pp; English.
XX
CC
     The invention provides membrane-bound PRO polypeptides and
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polynucleotides encoding them. The PRO sequences of the invention were identified based on extracellular domain homology screening. The PRO sequences have homology with proteins including LDL receptors, TIE ligands and various enzymes. The membrane-bound proteins and receptor molecules are useful as pharmaceutical and diagnostic agents. Receptor immunoadhesins, for instance, can be used as therapeutic agents to block receptor-ligand interactions. The membrane-bound proteins can also be employed for screening of potential peptide or small molecule inhibitors of the relevant receptor/ligand interaction. The PRO encoding sequences are useful as hybridization probes, in chromosome and gene mapping and in the generation of antisense RNA and DNA. PRO nucleic acid sequences will also be useful for the preparation of PRO polypeptides, especially by recombinant techniques.

XX SO Sequence 919 AA:

CC

Query Match 99.6%; Score 4760; DB 21; Length 919; Best Local Similarity 99.7%; Pred. No. 0: Matches 916: Conservative 0; Mismatches Indels 1; 2; Gaps 1; 1 MGLFRGFVFLLVLCLLHQSNTSFIKLNNNGFEDIVIVIDPSVPEDEKIIEQIEDMVTTAS 60 Qу 1 MGLFRGFVFLLVLCLLHQSNTSFIKLNNNGFEDIVIVIDPSVPEDEKIIEQIEDMVTTAS 60 Db 61 TYLFEATEKRFFFKNVSILIPENWKENPQYKRPKHENHKHADVIVAPPTLPGRDEPYTKQ 120 QУ 61 TYLFEATEKRFFFKNVSILIPENWKENPQYKRPKHENHKHADVIVAPPTLPGRDEPYTKQ 120 Db 121 FTECGEKGEYIHFTPDLLLEKKQNEYGPPGKLFVHEWAHLRWGVFDEYNEDQPFYRAKSK 180 Qу Db 121 FTECGEKGEYIHFTPDLLLGKKQNEYGPPGKLFVHEWAHLRWGVFDEYNEDOPFYRAKSK 180 181 KIEATRCSAGISGRNRVYKCQGGSCLSRACRIDSTTKLYGKDCQFFPDKVQTEKASIMFM 240 Qу 181 KIEATRCSAGISGRNRVYKCQGGSCLSRACRIDSTTKLYGKDCQFFPDKVQTEKASIMFM 240 Db Qy 241 QSIDSVVEFCNEKTHNQEAPSLQNIKCNFRSTWEVISNSEDFKNTIPMVTPPPPPPVFSLL 300 241 QSIDSVVEFCNEKTHNOEAPSLQNIKCNFRSTWEVISNSEDFKNTIPMVTPPPPPPVFSLL 300 Db Qу 301 KISQRIVCLVLDKSGSMGGKDRLNRMNQAAKHFLLQTVENGSWVGMVHFDSTATIVNKLI 360 301 KISQRIVCLVLDKSGSMGGKDRLNRMNQAAKHFLLQTVENGSWVGMVHFDSTATIVNKLI 360 Db 361 QIKSSDERNTLMAGLPTYPLGGTSICSGIKYAFQVIGELHSQLDGSEVLLLTDGEDNTAS 420 Qу 361 QIKSSDERNTLMAGLPTYPLGGTSICSGIKYAFQVIGELHSQLDGSEVLLLTDGEDNTAS 420 Db 421 SCIDEVKQSGAIVHFIALGRAADEAVIEMSKITGGSHFYVSDEAQNNGLIDAFGALTSGN 480 Qу 421 SCIDEVKOSGAIVHFIALGRAADEAVIEMSKITGGSHFYVSDEAQNNGLIDAFGALTSGN 480 Db 481 TDLSQKSLQLESKGLTLNSNAWMNDTVIIDSTVGKDTFFLITWNSLPPSISLWDPSGTIM 540 Qу 481 TDLSQKSLQLESKGLTLNSNAWMNDTVIIDSTVGKDTFFLITWNSLPPSISLWDPSGTIM 540 Db Qу 541 ENFTVDATSKMAYLSIPGTAKVGTWAYNLQAKANPETLTITVTSRAANSSVPPITVNAKM 600

Db	541		600
Qy	601	NKDVNSFPSPMIVYAEILQGYVPVLGANVTAFIESQNGHTEVLELLDNGAGADSFKNDGV	660
Db	601		660
Qy	661	YSRYFTAYTENGRYSLKVRAHGGANTARLKLRPPLNRAAYIPGWVVNGEIEANPPRPEID	720
Db	661	YSRYFTAYTENGRYSLKVRAHGGANTARLKLRPPLNRAAYIPGWVVNGEIEANPPRPEID	720
Qу	721	EDTQTTLEDFSRTASGGAFVVSQVPSLPLPDQYPPSQITDLDATVHEDKIILTWTAPGDN	780
Db	721	EDTQTTLEDFSRTASGGAFVVSQVPSLPLPDQYPPSQITDLDATVHEDKIILTWTAPGDN	780
Qу	781	FDVGKVQRYIIRISASILDLRDSFDDALQVNTTDLSPKEANSKESFAFKPENISEENATH	840
Db	781		840
Qу	841	IFIAIKSIDKSNLTSKVSNIAQVTLFIPQANPDDIDPTPTPTPTPDKSHNSGVNISTL	898
Db	841		900
Qy	899	VLSVIGSVVIVNFILSTTI 917	
Db	901		

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Fig. 258
RESULT 7
AAU29152
     AAU29152 standard; Protein; 919 AA.
XX
AC
     AAU29152;
XX
DT
     18-DEC-2001
                  (first entry)
XX
DE
     Human PRO polypeptide sequence (#129)
XX
KW
     PRO polypeptide; mammal; tumour; cancer; human; cattle; horse; sheep;
KW
     dog; cat; pig; goat; rabbit; tumour necrosis factor alpha; TNF-alpha;
KW
     blood; chondrocyte cell; cell proliferation; cell differentiation; colon;
KW
     adrenal; lung; breast; prostate; rectum; cervix; liver; genetic disorder.
XX
os
     Homo sapiens.
XX
PN
     WO200168848-A2.
XX
PD
     20-SEP-2001.
XX
     28-FEB-2001; 2001WO-US06520.
PF
XX
PR
     01-MAR-2000; 2000WO-US05601.
PR
     02-MAR-2000; 2000WO-US05841.
     03-MAR-2000; 2000US-187202P.
PR
     06-MAR-2000; 2000US-186968P.
PR
PR
     14-MAR-2000; 2000US-189320P.
PR
     14-MAR-2000; 2000US-189328P.
PR
     15-MAR-2000; 2000WO-US06884.
PR
     21-MAR-2000; 2000US-190828P.
PR
     21-MAR-2000; 2000US-191007P.
PR
     21-MAR-2000; 2000US-191048P.
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     21-MAR-2000; 2000US-191314P.
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     28-MAR-2000; 2000US-192655P.
PR
     29-MAR-2000; 2000US-193032P.
PR
     29-MAR-2000; 2000US-193053P.
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     30-MAR-2000; 2000WO-US08439.
PR
     04-APR-2000; 2000US-194449P.
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     04-APR-2000; 2000US-194647P.
PR
     11-APR-2000; 2000US-195975P.
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     11-APR-2000; 2000US-196000P.
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     11-APR-2000; 2000US-196187P.
PR
     11-APR-2000; 2000US-196690P.
PR
     11-APR-2000; 2000US-196820P.
PR
     18-APR-2000; 2000US-198121P.
PR
     18-APR-2000; 2000US-198585P.
PR
     25-APR-2000; 2000US-199397P.
     25-APR-2000; 2000US-199550P.
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     25-APR-2000; 2000US-199654P.
PR
     03-MAY-2000; 2000US-201516P.
PR
     17-MAY-2000; 2000WO-US13705.
PR
     22-MAY-2000; 2000WO-US14042.
PR
     30-MAY-2000; 2000WO-US14941:
PR
     02-JUN-2000; 2000WO-US15264.
PR
     05-JUN-2000; 2000US-209832P.
PR
     28-JUL-2000; 2000WO-US20710.
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22-AUG-2000; 2000US-0644848.
PR
PR
    24-AUG-2000; 2000WO-US23328.
PR
    08-NOV-2000; 2000WO-US30952.
PR
    01-DEC-2000; 2000WO-US32678.
    20-DEC-2000; 2000WO-US34956.
PR
XX
     (GETH ) GENENTECH INC.
PΑ
ХX
PΙ
    Baker KP, Chen J, Desnoyers L, Goddard A,
                                              Godowski PJ, Gurney AL;
PΙ
    Pan J, Smith V, Watanabe CK, Wood WI, Zhang Z;
XX
DR
    WPI; 2001-602746/68.
DR
    N-PSDB; AAS46053.
XX
PT
    Novel nucleic acids encoding PRO polypeptides, used to diagnose the
PT
    presence of tumours, such as prostate and breast tumours, in mammals and
    to screen for modulators of the compounds -
PT
XX
PS
    Claim 11; Fig 258; 774pp; English.
XX
CC
    Sequences AAU29024-AAU29328 represent PRO polypeptides of the invention.
    The PRO polypeptides and their associated nucleic acids can be used to
CC
CC
    detect the presence of a tumour in a mammal by comparing the level of
    expression of a PRO polypeptide in a test sample of cells from the animal
CC
CC
    and a control sample of normal cells, whereby a higher level of
CC
    expression in the test sample indicates the presence of a tumour in the
CC
    mammal. Mammals include dogs, cats, cattle, horses, sheep, pigs, goats
    and rabbits but are preferably human. The polypeptides can be used to
CC
CC
    stimulate tumour necrosis factor (TNF) alpha release from human blood,
    when contacted with it. A specific polypeptide can be used to stimulate
CC
CC
    the proliferation or differentiation of chondrocyte cells. The PRO
CC
    proteins can be used to determine the presence of tumours and also
CC
    susceptibility to tumour development, particularly adrenal, lung, colon,
    breast, prostate, rectal, cervical, or liver tumours, in mammalian
CC
CC
    subjects. The oligonucleotide probes specific for the PRO nucleic acids
CC
    can be used for genetic analysis of individuals with genetic disorders.
XX
SO
    Sequence
              919 AA;
 Query Match
                        99.6%; Score 4760; DB 22;
                                                  Length 919;
 Best Local Similarity
                       99.7%; Pred. No. 0;
 Matches 916; Conservative
                           0; Mismatches
                                              1;
                                                  Indels
                                                           2; Gaps
                                                                      1;
Qу
           1 MGLFRGFVFLLVLCLLHQSNTSFIKLNNNGFEDIVIVIDPSVPEDEKIIEQIEDMVTTAS 60
             Db
           1 MGLFRGFVFLLVLCLLHQSNTSFIKLNNNGFEDIVIVIDPSVPEDEKIIEQIEDMVTTAS 60
Qу
          61 TYLFEATEKRFFFKNVSILIPENWKENPQYKRPKHENHKHADVIVAPPTLPGRDEPYTKQ 120
             Db
          61 TYLFEATEKRFFFKNVSILIPENWKENPQYKRPKHENHKHADVIVAPPTLPGRDEPYTKQ 120
         121 FTECGEKGEYIHFTPDLLLEKKQNEYGPPGKLFVHEWAHLRWGVFDEYNEDQPFYRAKSK 180
Qу
             121 FTECGEKGEYIHFTPDLLLGKKQNEYGPPGKLFVHEWAHLRWGVFDEYNEDQPFYRAKSK 180
Db
Ov
         181 KIEATRCSAGISGRNRVYKCQGGSCLSRACRIDSTTKLYGKDCQFFPDKVQTEKASIMFM 240
```

מע	181	KIEATRCSAGISGRNRVYKCQGGSCLSRACRIDSTTKLYGKDCQFFPDKVQTEKASIMFM	240
Qy	241	QSIDSVVEFCNEKTHNQEAPSLQNIKCNFRSTWEVISNSEDFKNTIPMVTPPPPPPVFSLL	300
Db	241	QSIDSVVEFCNEKTHNQEAPSLQNIKCNFRSTWEVISNSEDFKNTIPMVTPPPPPVFSLL	300
Qy	301	KISQRIVCLVLDKSGSMGGKDRLNRMNQAAKHFLLQTVENGSWVGMVHFDSTATIVNKLI	360
Db	301	KISQRIVCLVLDKSGSMGGKDRLNRMNQAAKHFLLQTVENGSWVGMVHFDSTATIVNKLI	360
Qy	361	QIKSSDERNTLMAGLPTYPLGGTSICSGIKYAFQVIGELHSQLDGSEVLLLTDGEDNTAS	420
Db		QIKSSDERNTLMAGLPTYPLGGTSICSGIKYAFQVIGELHSQLDGSEVLLLTDGEDNTAS	
Qy		SCIDEVKQSGAIVHFIALGRAADEAVIEMSKITGGSHFYVSDEAQNNGLIDAFGALTSGN	
Db		SCIDEVKQSGAIVHFIALGRAADEAVIEMSKITGGSHFYVSDEAQNNGLIDAFGALTSGN	
Qy		TDLSQKSLQLESKGLTLNSNAWMNDTVIIDSTVGKDTFFLITWNSLPPSISLWDPSGTIM	
Db		TDLSQKSLQLESKGLTLNSNAWMNDTVIIDSTVGKDTFFLITWNSLPPSISLWDPSGTIM	
Qy		ENFTVDATSKMAYLSIPGTAKVGTWAYNLQAKANPETLTITVTSRAANSSVPPITVNAKM	
Db		ENFTVDATSKMAYLSIPGTAKVGTWAYNLQAKANPETLTITVTSRAANSSVPPITVNAKM	′
Qy		NKDVNSFPSPMIVYAEILQGYVPVLGANVTAFIESQNGHTEVLELLDNGAGADSFKNDGV	
Db		NKDVNSFPSPMIVYAEILQGYVPVLGANVTAFIESQNGHTEVLELLDNGAGADSFKNDGV	
Qy		YSRYFTAYTENGRYSLKVRAHGGANTARLKLRPPLNRAAYIPGWVVNGEIEANPPRPEID	720
Db ·		YSRYFTAYTENGRYSLKVRAHGGANTARLKLRPPLNRAAYIPGWVVNGEIEANPPRPEID	720
Qy Db		EDTQTTLEDFSRTASGGAFVVSQVPSLPLPDQYPPSQITDLDATVHEDKIILTWTAPGDN	780
		EDTQTTLEDFSRTASGGAFVVSQVPSLPLPDQYPPSQITDLDATVHEDKIILTWTAPGDN	
Qy Db		FDVGKVQRYIIRISASILDLRDSFDDALQVNTTDLSPKEANSKESFAFKPENISEENATH	
ДУ		FDVGKVQRYIIRISASILDLRDSFDDALQVNTTDLSPKEANSKESFAFKPENISEENATH	
Qy Db		IFIAIKSIDKSNLTSKVSNIAQVTLFIPQANPDDIDPTPTPTPTPDKSHNSGVNISTL	
Qу		IFIAIKSIDKSNLTSKVSNIAQVTLFIPQANPDDIDPTPTPTPTPTPDKSHNSGVNISTL VLSVIGSVVIVNFILSTTI 917	900
Qy Db			
עט	POT	ADOATOOAATAMLIDOIII ATA	